CLAIMS

- 1. Solid Lipid Nanoparticles of platinum compounds.
- 2. Solid Lipid Nanoparticles according to claim 1 wherein the platinum compounds are platinum complexes.
 - 3. Solid Lipid Nanoparticles according to claim 2, wherein the platinum complex is selected from trans-{bis[trans(diammine)(chloro)platinum (II)(\mu-1,6- hexanediamine)]}diammineplatinum tetranitrate salt of formula I

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Formula I

bis{trans(diammine)(chloro)platinum(II)} μ -(1,16-diamino-7,10-diazahexadecane-N1,N16) dinitrate salt. 2HNO₃ of formula II,

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Formula II

bis{trans(diammine)(chloro)platinum(II)} μ -(1,16-diamino-6,11-diazahexadecane-N1,N16) dinitrate salt. 2HNO₃ of formula III,

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Formula III

bis{trans(diammine)(chloro)platinum(II)}- μ -(1,12-diamino-4,9-diazadodecane-N₁,N₁₂) dinitrate salt. 2HNO₃ of formula IV,

Formula IV

bis{trans(diammine)(chloro)platinum

(II)}- μ -(1,8-diamino-4-azaoctane-

N¹,N⁸) dinitrate salt. HNO₃ of formula V,

Formula V

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- 4. A process for the preparation of Solid Lipid Nanoparticles of claims 1-3, comprising:
 - a) preparing a first microemulsion by mixing a molten lipid, a surfactant, and optionally a co-surfactant and the platinum compound acqueous solution;
 - b) preparing a solution by mixing a surfactant and optionally a co-surfactant in water, heating to complete solution, preferably at the same melting temperature of the lipid used in a) and adding a co-surfactant;

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- c) dispersing the microemulsion obtained in a) into the solution obtained in b) obtaining a multiple microemulsion c);
- d) dispersing the microemulsion obtained in c) in aqueous medium at a

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- temperature ranging from 0.5°C to 4°C obtaining a dispersion of solid lipid microspheres;
- e) washing with aqueous medium through ultrafiltration the obtained lipid microspheres obtained in d) and lyophilizing, optionally in the presence of a bulking agent and of a cryoprotecting agent.
- 5. Pharmaceutical compositions comprising the solid lipid nanoparticles of claims 1-3.
- A method of treating patients affected by cancer sensitive to platinum complexes which comprises administering to said patients a therapeutically
 effective amount of the solid lipid nanoparticles of claims 1-3.